

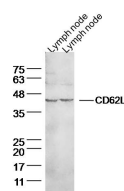
**bs-1036R****[ Primary Antibody ]****CD62L Rabbit pAb****Bioss**  
**ANTIBODIES**

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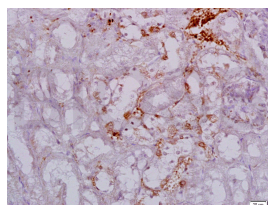
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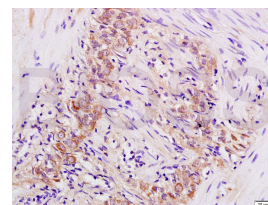
400-901-9800

**— DATASHEET —****Host:** Rabbit**Isotype:** IgG**Clonality:** Polyclonal**GeneID:** 20343**SWISS:** P18337**Target:** CD62L**Immunogen:** KLH conjugated synthetic peptide derived from mouse CD62L: 301-372/372. < Cytoplasmic >**Purification:** affinity purified by Protein A**Concentration:** 1mg/ml**Storage:** 0.01M TBS (pH7.4) with 1% BSA, 0.02% Proclin300 and 50% Glycerol.  
Shipped at 4°C. Store at -20°C for one year. Avoid repeated freeze/thaw cycles.**Background:** This gene encodes a cell surface adhesion molecule that belongs to a family of adhesion/homing receptors. The encoded protein contains a C-type lectin-like domain, a calcium-binding epidermal growth factor-like domain, and two short complement-like repeats. The gene product is required for binding and subsequent rolling of leucocytes on endothelial cells, facilitating their migration into secondary lymphoid organs and inflammation sites. Single-nucleotide polymorphisms in this gene have been associated with various diseases including immunoglobulin A nephropathy. Alternatively spliced transcript variants have been found for this gene. [provided by RefSeq, Oct 2009].**Applications:** **WB** (1:500-2000)**IHC-P** (1:100-500)**IHC-F** (1:100-500)**IF** (1:100-500)**Flow-Cyt** (1µg/Test)**Reactivity:** Mouse, Rat  
(predicted: Rabbit, Cow)**Predicted  
MW.:** 37 kDa**Subcellular  
Location:** Cell membrane**— VALIDATION IMAGES —**

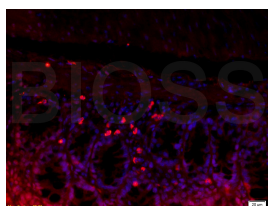
Sample: Lymph node (Mouse) Lysate at 40 µg  
Lymph node (Rat) Lysate at 40 µg  
Primary: Anti-CD62L (bs-1036R) at 1/300 dilution  
Secondary: IRDye800CW Goat Anti-Rabbit IgG at 1/20000 dilution  
Predicted band size: 37 kD  
Observed band size: 43 kD



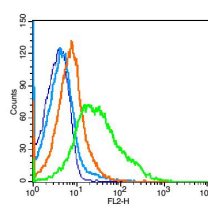
Tissue/cell: mouse kidney tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Incubation: Anti-L-Selectin Polyclonal Antibody, Unconjugated (bs-1036R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody (SP-0023) and DAB (C-0010) staining



Tissue/cell: rat colon tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (0.01M, pH 6.0), Boiling bathing for 15min; Block endogenous peroxidase by 3% Hydrogen peroxide for 30min; Blocking buffer (normal goat serum, C-0005) at 37°C for 20 min; Incubation: Anti-L-Selectin Polyclonal Antibody, Unconjugated (bs-1036R) 1:200, overnight at 4°C, followed by conjugation to the secondary antibody (SP-0023) and DAB (C-0010) staining



Tissue/cell: rat colitis tissue; 4% Paraformaldehyde-fixed and paraffin-embedded; Antigen retrieval: citrate buffer (



Blank control: mouse spleen cells (blue). Primary Antibody: Rabbit Anti-CD62L antibody (bs-1036R), Dilution: 1 µg in 100 µL 1X

**Important Note:** This product as supplied is intended for research use only, not for use in human, therapeutic or diagnostic applications.

0.01M, pH 6.0 ), Boiling bathing for 15min;  
Blocking buffer (normal goat serum,C-0005) at  
37°C for 20 min; Incubation: Anti-L-Selectin  
Polyclonal Antibody, Unconjugated(bs-1036R)  
1:200, overnight at 4°C; The secondary antibody  
was Goat Anti-Rabbit IgG, Cy3  
conjugated(bs-0295G-Cy3)used at 1:200 dilution  
for 40 minutes at 37°C. DAPI(5ug/ml,blue,C-0033)  
was used to stain the cell nuclei

PBS containing 0.5% BSA; Isotype Control  
Antibody: Rabbit IgG(orange) ,used under the  
same conditions ); Secondary Antibody: Goat  
anti-rabbit IgG-PE(white blue), Dilution: 1:200 in  
1 X PBS containing 0.5% BSA. Protocol The cells  
were fixed with 2% paraformaldehyde (10 min).  
Primary antibody (bs-1036R, 1μg /1x10<sup>6</sup> cells)  
were incubated for 30 min on the ice, followed  
by 1 X PBS containing 0.5% BSA + 1 0% goat  
serum (15 min) to block non-specific protein-  
protein interactions. Then the Goat Anti-rabbit  
IgG/PE antibody was added into the blocking  
buffer mentioned above to react with the  
primary antibody at 1/200 dilution for 30 min on  
ice. Acquisition of 20,000 events was performed.

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## — SELECTED CITATIONS —

- **[IF=7.3]** Ye Yuqing. et al. Dynamic changes of immunocyte subpopulations in thermogenic activation of adipose tissues. FRONT IMMUNOL. 2024 May;15: IF ;Mouse. 38812501